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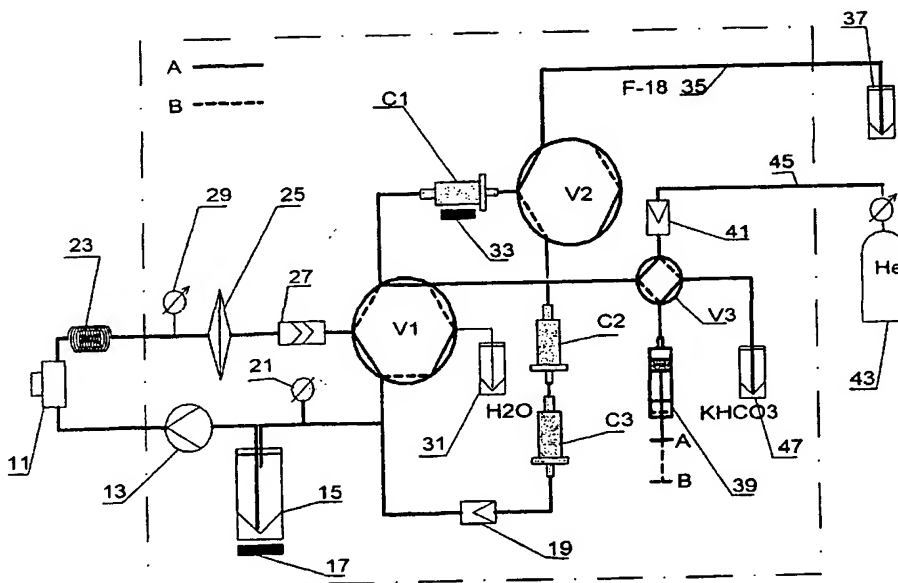
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(54) Title: PROCESS AND APPARATUS FOR PRODUCTION OF F-18 FLUORIDE



**(57) Abstract:** A process and apparatus for producing the  $^{18}\text{F}$  isotope from water enriched with the  $^{18}\text{O}$  isotope using high energy protons a cyclotron. The apparatus has a cyclotron target cavity (11) that is connected to a target loop (L1) that contains a water reservoir (15), pump (13), and pressure regulator (27). Water is continuously recirculated through the target cavity to increase re-liability. After irradiation long enough to produce a desired amount of  $^{18}\text{F}$ , water in the target loop (L1) is diverted through an  $^{18}\text{F}$  extraction device (C1) before being returned to the target loop. Preferably, the returning water is purified and additional water may be added to the target loop as needed to permit continuous irradiation and production of  $^{18}\text{F}$ .

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*For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.*

## PROCESS AND APPARATUS FOR PRODUCTION OF F-18 FLUORIDE

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## BACKGROUND

## 1. Technical field.

10 The invention relates to production of an  $^{18}\text{F}$  radioisotope by means of proton irradiation of  $^{18}\text{O}$  enriched water.

## 2. Background.

15 The  $^{18}\text{F}$  isotope (hereinafter, F-18 isotope or F-18) has become widely used in nuclear medicine for diagnostic studies using a Positron Emission Tomography (PET) body scanning technique. The F-18 is typically used to label an injectable glucose derivative. Because of its short half-life (109 min.), this isotope must be used as soon as possible after production. This makes it impossible to accumulate a sufficient quantity for delayed use. Therefore, work shifts usually start near midnight with production for distant (via automobile) hospitals first, followed by that for  
20 nearby hospitals in the very early morning. Any shortage in production has an immediate and direct effect on users. As a result, reliability and predictability of production are extremely important for users as well as suppliers of this isotope.

The two main methods of producing F-18 use an  $^{18}\text{O}(\text{p},\text{n})^{18}\text{F}$  reaction in a cyclotron. Both gaseous oxygen and liquid water enriched with  $^{18}\text{O}$  (hereinafter, O-18) have been used as target  
25 materials. However, the gaseous approach is very difficult in practice because the F-18 is very reactive and hard to recover from a gaseous medium. The overwhelming majority of production facilities use water enriched with O-18 ( $\text{H}_2[^{18}\text{O}]$ , hereinafter, O-18 water).

Using O-18 water is not without problems, also. For production efficiency, it is desirable to use water that is as much enriched as possible. However, 95% enriched O-18 water costs  
30 approximately \$150 per ml. Also, PET has been gaining greater acceptance and the building of new O-18 water production facilities is lagging behind demand. The cost pressures make conservation and reuse of the O-18 water target material even more important.

In a typical system for F-18 production, the target is typically loaded with a pre-determined amount of O-18 water by means of a syringe or pump. The volume of water in the target is about 0.8 ml, but another 1-2 ml is required to fill the lines leading to the target. The water delivery system is then isolated from the target by means of a valve and the target is irradiated. This can be described as a "static" target, meaning that the target material remains in the target throughout the irradiation time.

The irradiated water is then removed from the target, typically by means of inert gas pressure, and transported over a delivery line leading outside the cyclotron shielding to a collection vial about 8 m (25 feet) from the target. The F-18 isotope is then separated from the water and processed for production of a radiopharmaceutical agent.

A considerable amount of O-18, typically 25-30%, is lost after each run. The O-18 isotope is used up in three ways. First, a very small amount, on the order of nanoliters, is actually converted to F-18. The next most important loss of O-18 is due to a combination of leakage and isotopic exchange with  $^{16}\text{O}$  oxides in the target, transport lines and storage vessels. After one run of an hour or two, the enrichment factor can drop from 95% to 85-90%. This is still high enough to be economical to run a cyclotron, but the amount of contamination is too high, as will be explained below. (As the enrichment factor falls, the irradiation time increases. 80% is a minimum under current economic conditions.)

The third loss is due to leakage of target material from the pressurized target and attached tubing which may lead to a reduced water level in the target and, if severe enough, to a catastrophic failure. Target cooling relies on the liquid water material present in the target to function as a heat conductor. A typical 1 ml target must dissipate over 500W of heat for as long as 2-3 hours. Many target systems are pressurized to as high as 3.4 MPa (500 psig) or higher to improve target thermal stability. In these conditions, containment of a small amount of water becomes a significant technical problem. Loss of a very small amount of target material may have dramatic consequences such as target foil rupture, target body degradation, and loss of target yield.

Although 70-75% of the initial O-18 water remains, the biggest effective loss is due to contamination. Any contamination in the liquid water increases the formation of super-heated steam with increased leakage and loss of cooling. Because the consequences are so adverse, the water recovered after only one run in a static target system must be sent back to the supplier for reprocessing to remove contaminants.

Existing static target systems do not provide any mechanism to timely detect the critical loss of target material during irradiation. In addition, in a static target it is impossible to monitor the amount of radioactive F-18 being produced with any certainty. The result of a production run may not be known until after its completion, up to several hours after start of production. Given the fact that production and delivery schedules do not allow much flexibility due to the extremely short half-life of the F-18, this uncertainty results in a decrease in reliability and availability of the product.

## INVENTION SUMMARY

Accordingly, one objective of the invention is to increase the reliability of the production of F-18 from O-18 enriched water irradiated by high energy protons produced by a cyclotron. Further objectives are to increase the efficiency so that the cyclotron can be irradiating O-18 without interruption. Still another objective is to continually reuse O-18 water from which F-18 is periodically extracted. Another objective is to be able to add additional new O-18 water as it is lost due to system leakage and the like so that the system can run for an extended period without interruption.

These objectives and more are realized with a process that continuously recirculates O-18 enriched water through a target loop that includes a target cavity for a cyclotron that irradiates the target cavity with protons to convert a portion of O-18 to F-18.

Longer irradiation without failure is achieved by using a combination of one or more of the following: maintaining a pressure of at least about 1.7 MPa (250 psig) in the target cavity; recirculating the O-18 water through the target cavity at least about once every two minutes; and maintaining an O-18 water volume in the target loop that is at least about ten times the volume of the target cavity, itself. Additional benefit can be obtained by substantially cooling the O-18 water after exiting the target cavity and before reintroduction.

Increased efficiency is obtained by periodically recharging the target loop with additional O-18 water without interrupting irradiation and using protons having an energy of about 16 Mev and an intensity of at least about 40  $\mu$ A on the target cavity.

Rather than stop irradiation and loose cyclotron time, F-18 can be extracted from irradiated O-18 water in the target loop by periodically, e.g., every hour or two, briefly diverting the target loop through an F-18 extraction device without interrupting irradiation of the target cavity.

Because the amount of O-18 that is converted to F-18 is quite small, e.g., less than 0.1% of the O-18 is converted, after F-18 is extracted, the remaining O-18 water can be purified by solid phase purification devices and reintroduced into the target loop.

The aforementioned target loop can be implemented with, in order: an O-18 water reservoir; a pump; a target cavity; and a backpressure regulator. The pump must be capable of generating the minimum desirable pressures of 1.7 MPa (250 psig) and, for a typical target loop volume of 10 ml, a flow rate of 2 ml/min. Cooling of the O-18 water may be accomplished with a coil of tubing connected on the output side of the target cavity.

The F-18 may be recovered from some types of F-18 extraction devices, e.g., anion exchange types, with an eluant and a gas source for forcing the F-18 eluate into a delivery vial.

O-18 water purification devices are preferably connected through a valve to the output of the F-18 extraction device and may reintroduce O-18 water into the target loop by means of a simple check valve.

Production efficiency can be further increased by having a source vial with new O-18 water to periodically, without stopping irradiation, recharge the target loop as O-18 water is used up due to leakage and the like.

Valves and tubes are provided to controllably connect various elements to perform various functions to carry out the invention.

## BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 is a schematic diagram of apparatus for practicing the invention;

FIG. 2 is a graph of the reservoir vial and exchange cartridge radioactivity for two experimental runs;

FIG. 3 is a graph of target water conductivity for the same runs as in FIG 2; and

FIG. 4 is a graph of target water pressure for the same runs as in FIG. 2.

## DETAILED DESCRIPTION OF MODES FOR CARRYING OUT THE INVENTION

FIG. 1 is a schematic diagram of the apparatus whose component parts will now be described. All of these are used in the field of High Pressure Liquid Chromatography (HPLC) where they are fairly common. Connections between components were made with either 1/16 in. (1.6 mm) OD type 316 stainless steel tubing or 1/16 in. (1.6 mm) OD, 0.030 in. (0.8 mm) ID polyetheretherketone (PEEK) tubing, as was mechanically convenient. The choice of tubing is

believed to be not critical. PEEK compression fittings are used for both types of tubing.

The target (11) is the standard "high yield" cyclotron target supplied by General Electric (U.S.) PET Systems AB (Uppsala, Sweden). This target has a silver body with an 0.8 ml target volume behind a 1 cm diameter circular aperture covered with a cobalt alloy Havar (TM) (Co 42.5%, Cr 20%, Ni 13%, Fe/W/Mo/Mn) foil sealed with a crushed silver o-ring. Using standard components (not illustrated), the target body is cooled by 20 °C water and the aperture foil is cooled with 340 kPa (50 psig) room temperature helium gas. Use of PEEK fittings means that the target is electrically insulated from the remainder of the apparatus. Thus, the beam current absorbed by the target material can be measured with an ammeter (not shown) connected between the target (11) and the cyclotron ground.

The cyclotron used is a standard one from the target supplier and is not illustrated. It is a model PETtrace (TM) 2000 negative ion type that accelerates singly negatively charged hydrogen ions. The cyclotron produces a close to Gaussian beam of 16.5 MeV protons with a total beam current of up to 75 uA. As is usual, tungsten collimators are used to center a more uniform beam distribution in the 1 cm diameter target aperture. A carbon foil in the cyclotron beam strips electrons from the negatively charged hydrogen ions to produce protons (positively charged hydrogen ions).

The input to the target is supplied with O-18 water by a pump (13) that is in turn connected to a reservoir vial (15) with a capacity of about 5 ml. The pump is a Cole Palmer (Vernon Hills, IL) model U-07143-86 single piston type. This pump has a sapphire piston, ruby valve seats, gold-plated stainless steel springs, and type 317 stainless steel housings and fittings. Other wetted parts are made from non-reactive materials such as PEEK. The flow rate is set to about 5 ml/min.

A reservoir vial radiation sensor (17) is used to monitor radiation in the vial (15). This sensor is constructed with a 5 mm NaI scintillation crystal epoxied to a photodiode. (A PMT is not needed.) The assembly is within 1.25 cm (1/2 in.) of the vial (15), but a photocurrent amplifier (not illustrated) is located 3 m (10 feet) away to reduce the effects of a neutron flux generated by the irradiated target.

The input to the vial (15) comes from a valve (V1) in parallel with an Upchurch (Oak Harbor, WA) model CV-3302 liquid check valve (19). This line is also connected to a Cole Palmer digital conductivity meter (21) having a micro-flow cell consisting of a 1.6 mm (1/16 in.) ID glass tube with embedded platinum electrodes.

Valve (V1) is a Rheodyne (Rohnert Park, CA) model 7000 pneumatically actuated 6-port

with two positions, A and B, indicated by the solid and dashed lines, respectively. In position A, 3 pairs of adjacent ports are connected, while in position B, the three other adjacent pairs are connected. As illustrated, one of the ports is sealed off. The pneumatic actuator gas lines are not illustrated.

5 The output of the target (11) goes through a cooling coil (23) that consists of 3-m (10 feet) of loose 5 cm (2 in.) dia coils of 1.6 mm (1/16 in.) OD stainless tubing. The cooling coil is essentially suspended in ambient air and provides cooling for water exiting the target (11). The coil is connected to an Alltech (Deerfield, IL) 10 micron stainless steel filter (25) that filters out, e.g., silver particles, that may have been picked up in the target. The filter is connected to an  
10 Upchurch model U-469 back pressure regulator (27) adjustable in the range of 1.7 - 3.4 MPa (250 - 500 psig). The pressure in the volume after the pump (13) is monitored by an Omega Engineering (Stamford, CT) model PX176-500 0 - 3.4 MPa (0-500 psig) pressure transducer (29). It is well know that higher pressures in the target volume increases the boiling point allowing higher intensity irradiation. However the present apparatus leaked at 3.4 MPa (500 psig) and the  
15 maximum pressure could not be used.

When valve (V1) is in the A position, the pump (13) circulates water through the target loop (L1). Circulation is at the rate of about 5 ml/min. With a calculated loop volume of about 5 ml added to the reservoir vial (15) volume of 5 ml to yield 10 ml, this means that 2 minutes is required for one round trip.

20 The initial source of O-18 water is source vial (31) that is connected to one of the ports of valve (V1). This vial has a 50 ml capacity. The concentration of the O-18 isotope is not necessarily 100%. Any concentration can be used, but in normal production, at least 80% and preferably higher should be used to reduce irradiation time and the cost of the cyclotron.

A Waters (Franklin, MA) model SepPak (TM) QMA cartridge (C1) containing silica  
25 derivatized by quaternary ammonia is connected between the valve (V1) and a second valve (V2). This cartridge can adsorb F-18 ions from water. The F-18 can then be extracted using eluants such as 20-40 mM sodium or potassium carbonate in water or a water/acetonitrile mixture. The amount of F-18 in cartridge (C1) is monitored by the photodiode sensor (33) adjacent to the cartridge.

30 Valve (V2) is also a Rheodyne series 7000 pneumatically actuated 6-port with positions A and B as indicated by the solid and dashed line, respectively. Only half of this valve is used. One side of valve (V2) is connected to an F-18 delivery line (35) constructed from 1.6 mm (1/16



in.) OD PEEK tubing stretching about 8 m (25 feet) from the cyclotron target area to an F-18 delivery vial (37).

The other side of valve (V2) is connected to an in-line pair of deionizing cartridges (C2) and (C3) that are connected to the check valve (19). These are used to remove impurities from the O-18 water, especially in later stages of a production run. Cartridge (C2) is an Alltech (Deerfield, IL) MaxiClean (TM) model SCX (Strong Cation Exchange) cartridge containing 600 mg of polystyrene resin derivatized with sulfonic acid. Cartridge (C3) is a similar model SAX (Strong Anion Exchange) cartridge derivatized with a tetra-alkylammonium compound. Check valve (19) prevents back flow into these cartridges.

A third valve (V3) is connected to valve (V1). This is a model HVP-E 86779 4-port supplied by Alltech. One of these ports is connected to a Hamilton Gastight (TM) model 1002 2.5 ml syringe pump (39) (supplied by Alltech) with a pneumatically actuated plunger. The pump body is glass while the plunger is made from polytetrafluorethylene with the trade name Teflon. As shown, the plunger has two extreme positions, all the way in, designated A, and all the way out, designated B.

Another port of valve (V3) is connected to a gas check valve (41) that is connected to a remote helium tank (43) via helium line (45). The tank is filled with Matheson UHP grade 5.5 (i.e. 99.9995% pure) helium. The other port of valve (V3) is connected to an eluant vial (47) containing a suitable eluant solution such as a sodium carbonate solution in water.

All components shown inside the large box in FIG. 1 are mounted on and between two 20 cm (8 in.) wide by 36 cm (14 in.) high by 6 mm (1/4 in.) thick aluminum plates separated by 15 cm (6 in.). This is about the same volume used by the standard liquid target filler apparatus supplied by the cyclotron manufacturer. This assembly is placed within 60-90 cm (2 - 3 feet) of the target (11). In addition to F-18 delivery line (35) and helium line (45), all other pneumatic actuator and electrical lines are brought outside the cyclotron radiation shield. While it would reduce the number of long lines to bring all components except the target loop (L1) outside the shield, this would require a long line to the O-18 source vial (31) that would increase the possibility of contaminating the O-18 water.

The apparatus is operated under control of an IBM PC compatible computer and control system (not illustrated) based on an Omega Engineering (Stamford, CN) model CIO DAS 08 I/O board having analog and digital input and digital output ports. The output ports drive local solenoids that, in turn, drive pneumatic actuators located with the apparatus. In order to monitor

operation, the computer also stores in memory readings from the pressure, radiation, and conductivity meters.

#### Operation:

As noted above, production of F-18 for medical uses takes place in a work shift just preceding the beginning of a hospital day. Operation of the apparatus illustrated in FIG. 1 can be carried out with a series of runs that would typically last an hour or more. Before a run starts, it is necessary to make sure that the target loop (L1) is filled with O-18 water. Then, a second production sequence of steps would produce F-18, extract the F-18 produced, and deliver it to the external vial (37) for further processing.

When the system is first assembled, the first requirement is to fill the target (11) and reservoir vial (15) with O-18 water. This is accomplished by connecting the vial of O-18 water (31) to valve (V1). The three valves in the system and the syringe pump (39) are sequenced according to the following Table I.

Table I: Fill Target Loop Sequence

Step	(V1)	(V2)	(V3)	Syringe	Typical Time (s)
1. Start	A	A	A	A (in)	-
2. Fill Syringe	B	A	B	B (out)	5
3. Switch Valves	A	B	B	B (out)	5
4. Add Water	A	B	B	A (in)	10
5. Purge Cartridges	A	B	A	A (in)	5
6. Reset Valves	A	A	A	A (in)	2

In the fill syringe step, O-18 vial (31) is connected through valves (V1) and (V2) to syringe (39). Then, when the syringe plunger is pulled out, O-18 water is pulled from the vial into the syringe.

In the switch valves step, the syringe is connected through valve (V1) to cartridge (C1) and through valve (V2) to cartridges (C2) and (C3). In the add water step, the plunger of syringe (39) is pushed in and O-18 water is forced through the cartridges (C1), (C2), and (C3) and check valve (19) into the reservoir vial (15). The volume and stroke of syringe (39) was adjusted to produce an injection of about 0.75 ml. The volume of the cartridges and connecting lines is about 1-2 ml.

This particular arrangement means that the initial charge of reservoir vial (15) as well as any subsequent recharges with O-18 water will be purified by the ion exchange cartridges (C2) and (C3). In the purge cartridges step, Valve (V3) connects the 340 kPa (50 psig) helium supply (43) via valve (V1) to cartridge (C1) and via valve (V2) to cartridges (C2), and (C3). This purges the cartridges and forces any remaining water into reservoir vial (15). In the reset valves step, valve (V2) is returned to the A position disconnecting cartridge (C1) from cartridges (C2) and (C3), in preparation for either a repeat of the fill target sequence or the production sequence.

When a system is first assembled, the fill target sequence is repeated about 15 times to fill the loop (L1), containing the target (11) and reservoir vial (15), with total of 10 ml of water. In the beginning of a work shift, the fill target sequence is repeated as necessary until reservoir vial (15) contains about 5 ml of water. After completion of the fill target sequences at the beginning of a work shift, the pump (13) and the cyclotron are turned on and left on for the remainder of the shift. Next is a production sequence of steps as listed in Table II.

Table II: Production Sequence:

Step	(V1)	(V2)	(V3)	Syringe	Typical Time (s)
1. Irradiation	A	A	B	A (in)	300 and up
2. Extraction	B	B	A	A (in)	360
3. Purge	A	B	A	A (in)	20
4. Fill Syringe	A	B	A	B (out)	10
5. Prepare to Deliver	A	A	B	B (out)	2
6. Elute F-18	A	A	B	A (in)	15
7. Deliver F-18	A	A	A	A (in)	240
8. Reset valves	A	A	B	A (in)	1

During the Irradiation step, the cyclotron is turned on and the target (11) is irradiated. With valve (V1) in the A position, pump (13) is running and circulates water through the target loop (L1). Check valve (19) blocks circulation back into the cartridges (C2) and (C3). Back-pressure regulator (27) maintains the pressure at some level between 1.7 – 3.4 MPa (250 - 500 psig). Pressure monitor (29), that is upstream of the 10-micron filter (25), signals the control system if an over or under-pressure occurs. The conductivity monitor (21) signals the control system if the

conductivity is too high, indicating excessive contamination. During irradiation, the amount of F-18 created is monitored by the reservoir vial radiation sensor (17) and associated circuitry.

With valve (V3) in the B position, the helium supply pressurizes the eluant vial (47), but has no other effect. With valve (V2) and the syringe (39) in the A position, there is no flow  
5 through the cartridge (C1).

After a desired amount of F-18 has accumulated in the target, it is extracted. Valves (V1) and (V2) are switched to the B position breaking the loop ((L1)) at valve (V1) and forming a loop through the cartridges (C1), (C2), and (C3). QMA cartridge (C1) retains F-18 while deionizing cartridges (C2) and (C3) remove impurities from the water. After 360 sec., about 85%-90% of the  
10 F-18 has been absorbed on the cartridge.

The F-18 level in the QMA cartridge (C1) is monitored by the photodiode (33) and the conductivity of the water is monitored by the photodiode (17).

In the purge step, as much O-18 water as possible is removed from the QMA cartridge (C1). Valve (V1) is switched to the A position connecting the cartridge through valve (V3) to the  
15 helium source (43) and reestablishing the target loop (L1). The helium gas pressure pushes water from the QMA cartridges through the deionizing cartridges (C2) and (C3) and past the check valve (19) into vial (15).

The next four steps deliver F-18 to the delivery vial (37). With valve (V3) in the A position, the syringe (39) is connected to the eluant vial (47). Pulling the plunger out fills the  
20 syringe with about 0.75 ml of eluant. This takes about 10 seconds. Then, valve (V2) is switched to the A position and valve (V3) to the B position. This connects the syringe (39) to the QMA cartridge (C1) and from there to the delivery vial (37). In the elute step, the plunger of the syringe (39) is pushed in over about a 15 second period. This forces eluant solution into the QMA cartridge (C1).

25 Next, in the delivery step, valve (C3) is switched to the A position so that the helium source (43) is connected to the QMA cartridge (C1). The helium gas pressure forces the F-18 containing eluate into the delivery tube (35) and to the delivery vial (37). This takes about 240 seconds.

The recovery steps, starting with filling the syringe (39) and ending with delivery, are then  
30 repeated to accomplish complete removal of F-18 from the cartridge (C1). About 85% of F-18 produced in the target (11) is removed from the QMA cartridge (C1) after two extractions. This estimate is based on known target production efficiency as compared to the amount of F-18

delivered into the receiving vial (37).

A fraction of the remaining 15% of F-18 will be recovered in a subsequent production sequence depending on the length of the next run compared to the 109 minute F-18 half-life.

At the conclusion, valve (V3) is switched back to position B to begin another production sequence or left in position A if the target-loop (L1) needs replenishing with water using the Fill Target sequence.

#### Four Working Examples:

Four consecutive trial runs were made without shutting down the system using the same set of cartridges. Two sets of beam current amounts and irradiation times were used. The concentration of O-18 in the starting water was only 80% (because of the expense of higher concentrations). The eluant was 40 mM sodium carbonate solution in water. A Capintec (Ramsey, NJ) 7BT dose calibrator was used to measure the amount of recovered F-18 after each run. The results appear in Table III.

Table III: Four Trial Runs:

<u>Run #:</u>	<u>Beam Current (uA)</u>	<u>Irradiation Time (min)</u>	<u>Recovered F-18 (mCi)</u>
1	20	5	98
2	20	5	91
3	40	126	2240
4	40	104	2730

Runs 1 and 2 are too short to produce useful amounts of F-18, but were truncated to check system operation. In principal, the F-18 from many short runs can be combined, but this produces a very dilute solution of F-18. Therefore, a continuous run that delivers 2 - 4 Ci is preferred.

The higher amount of F18 delivered in run 4, despite a shorter irradiation time, is due to activity remaining in the target loop (L1), including the target (11) and the reservoir vial (15), after run 3. There also were two extraction steps performed in run 4 as compared to one extraction step in run 3 which leads to a more complete extraction of the isotope. Further, it is not unusual with prior art static systems for recoveries to vary by 5-10% between otherwise identical runs.

For runs 3 and 4, FIG. 2 shows the radioactivity in the reservoir vial (15) and QMA cartridge (C1) as determined by sensors (17) and (33), respectively, as a function of time, T, in

hours and minutes. The output of these two sensors were scaled to approximate the recovered F-18. The only steps that are long enough to see on this scale of hours and minutes are irradiation, extraction and delivery.

At the beginning of run 3, radioactivity in the reservoir vial 15, indicated by the solid trace, builds up approximately exponentially because the irradiation time is comparable to F-18's 1 hour and 49 minute half-life. At approximately 2:28, extraction starts and the amount of F-18 in the reservoir vial (15) drops rapidly with a corresponding increase in the QMA cartridge (C1) indicated by the dotted line. The irradiation continues during the extraction step which is why the F-18 amount is still rising when the elution step starts at approximately 2:38. This leaves some F-18, some of which is produced during extraction of run 3, in the reservoir vial (15) at the start of irradiation run 4.

Although not visible in the graphs because the Fill Target Loop Sequence takes less than 30 seconds, at the end of run 3, the target loop (L1) was recharged with approximately 1.5 ml of O-18 water. This particular target was used for these experiments because it leaked too much to be used in a normal static target production run. The 1.5 ml added was an estimate based on a prior leak test without irradiation. The basic requirement is that the target (11) not run dry. This is fulfilled, if the out take tube of the reservoir vial (15) is always submerged. This is not difficult because, through experience, an estimate can be made of target loop water losses and the Fill Target Loop Sequence can be performed at any time as needed.

At approximately 4:19, at the end of radiation run 4, F-18 is extracted, but with greater apparent efficiency than after run 3. This is followed by a short delivery step and then a second extraction step ending just before the graph. What would have been Run 5 was terminated because cyclotron time allocated to the experiments ran out. It is believed that runs could have continued until the O-18 water in the source vial (31) ran out.

FIG. 3 shows the target loop (L1) water conductivity over the same time period as in FIG. 2. This increases with time due to the buildup of various ionic species produced mainly by target corrosion and decreases due to the SAX and SCX cartridges (C2) and (C3) during the extraction step. (Note that, F-18 does not contribute to conductivity changes because it is not present in chemically significant quantities.) The fact that the conductivity returns back to low levels after isotope extraction demonstrates the possibility of indefinite reuse of target material contained in the loop (L1) and reservoir (15).

FIG. 4 shows the pressure in the target (11) over the same time period as in FIG. 2. It is held relatively constant by the pressure regulator with an increase when the target loop is diverted through the cartridges during extraction steps.

5 Alternative approaches:

The above example of operation and system description are provided to illustrate one of many ways to accomplish the recirculation and extraction. A variety of similar components may be used with equal success. For example, any high-pressure piston pump designed for HPLC or similar application and equipped with inert piston and check valves can be used to pump liquid. Similarly, a variety of valve designs are available that could be used to substitute Hamilton and Rheodyne valves provided that they utilize inert materials and are capable of sustaining the required pressure and are compatible with water.

Plumbing of the system can be substituted with all stainless steel or plastic material. Appropriate materials can be used to replace PEEK or type 316 stainless steel. Additional cooling of water removed from the target by means of a heat exchanger may be beneficial. Additional pressure, radioactivity and temperature sensors could provide better feedback and monitoring.

It may be beneficial to use increased water flow rates to provide better mixing inside the target and to achieve better heat dissipation. With higher water flow rates and additional cooling it may be possible to significantly increase beam current deposited into the target, thus increasing the isotope production rate. Thus, a recirculating target design has the potential to significantly increase production of the isotope.

The single syringe was a convenient device for transferring O-18 water and eluant. However, with a different valve arrangement, two syringes could be used or different fluid transfer devices substituted. For example, gas pressure could be used to force fluids out of containers.

A wide variety of commercially available cartridges designed for solid phase extraction and ion exchange can be used to substitute for QMA, SAX or SCX cartridges. Additional cartridges and filters can be installed as necessary to remove other potentially harmful impurities, such as a type C-18 cartridge to remove organic materials. Additionally, a sterilizing filter can be incorporated in a purification loop to remove microbial contamination, if necessary.

Various solutions can be used to remove extracted F-18 isotope from the QMA cartridge to accommodate requirements of the chemical processing that follow isotope production, as long as these solutions have sufficient ion strength to equilibrate the QMA cartridge and displace fluoride

ion. For example, a solution of a tetraalkylammonium base or salt such as tetrabutyl ammonium carbonate or potassium carbonate in an equimolar mixture with a polycyclic aminopolyether such as 4,7,13,16,21,24-hexaoxa-1,10-diazabicyclo[8,8,8]hexacosane can be used to provide increased reactivity of F-18 fluoride in following nucleophilic substitution reactions. Such a solution can be  
5 used directly in the synthesis of some useful radiopharmaceutical agents such as [F18]  
2-Deoxy-2-Fluoro-D-glucose, thus eliminating one step from the synthesis procedure and increasing yield and reducing synthesis time.

Lastly, the invention is not limited to using the particular target and cyclotron employed for the trial runs. Equivalents from other manufacturers should require only minor changes in  
10 apparatus.

It should therefore be clear that the detailed description of one working embodiment does not prevent inclusion of other equivalent embodiments within the purview of the invention that is defined by the following claims.



What is claimed is:

1. A process of making an F-18 isotope by using a proton beam to irradiate a target containing recirculating O-18 water characterized in that the O-18 water is also circulated through a loop containing an F-18 extraction device and then back to the target.

2. The process of claim 1 wherein said F-18 extraction device is an anion exchange type filter.

3. The process of claim 1 wherein the pressure in the re-circulating O-18 water in the target is maintained at not less than 1.7 MPa.

4. The process of claim 1 further including the step of purifying the O-18 water with solid phase purification devices before re-circulation to the target cavity.

5. The process of claim 1 further including the step of periodically recharging the target with additional O-18 water without interrupting the proton irradiation.

6. The process of any of claims 2, 3, 4, or 5 wherein the F-18 extraction is performed periodically without significantly interrupting proton irradiation.

7. An apparatus for making an F-18 isotope that uses a proton beam to irradiate a target containing re-circulating O-18 water characterized by the addition of an F-18 extraction device introduced in the path of the recirculating O-18.

8. The apparatus of claim 7 wherein the F-18 extraction device is an anion exchange type filter.

9. The apparatus of claim 7 wherein a backpressure regulator is used to maintain the O-18 water pressure.

10. The apparatus of claim 7 wherein solid phase purification cartridges are used to purify the O-18 water before recirculation to the target cavity.

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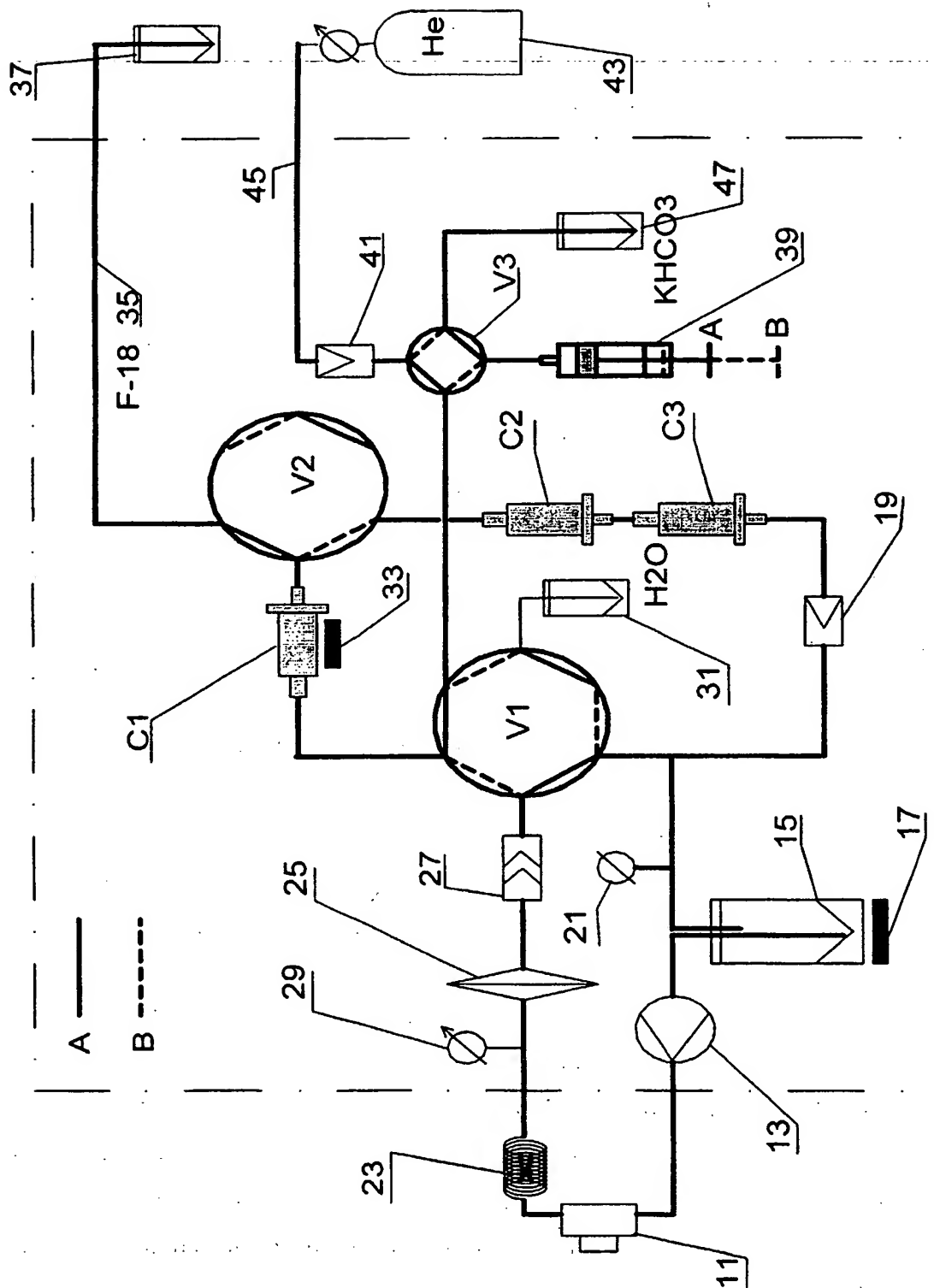


FIG. 1

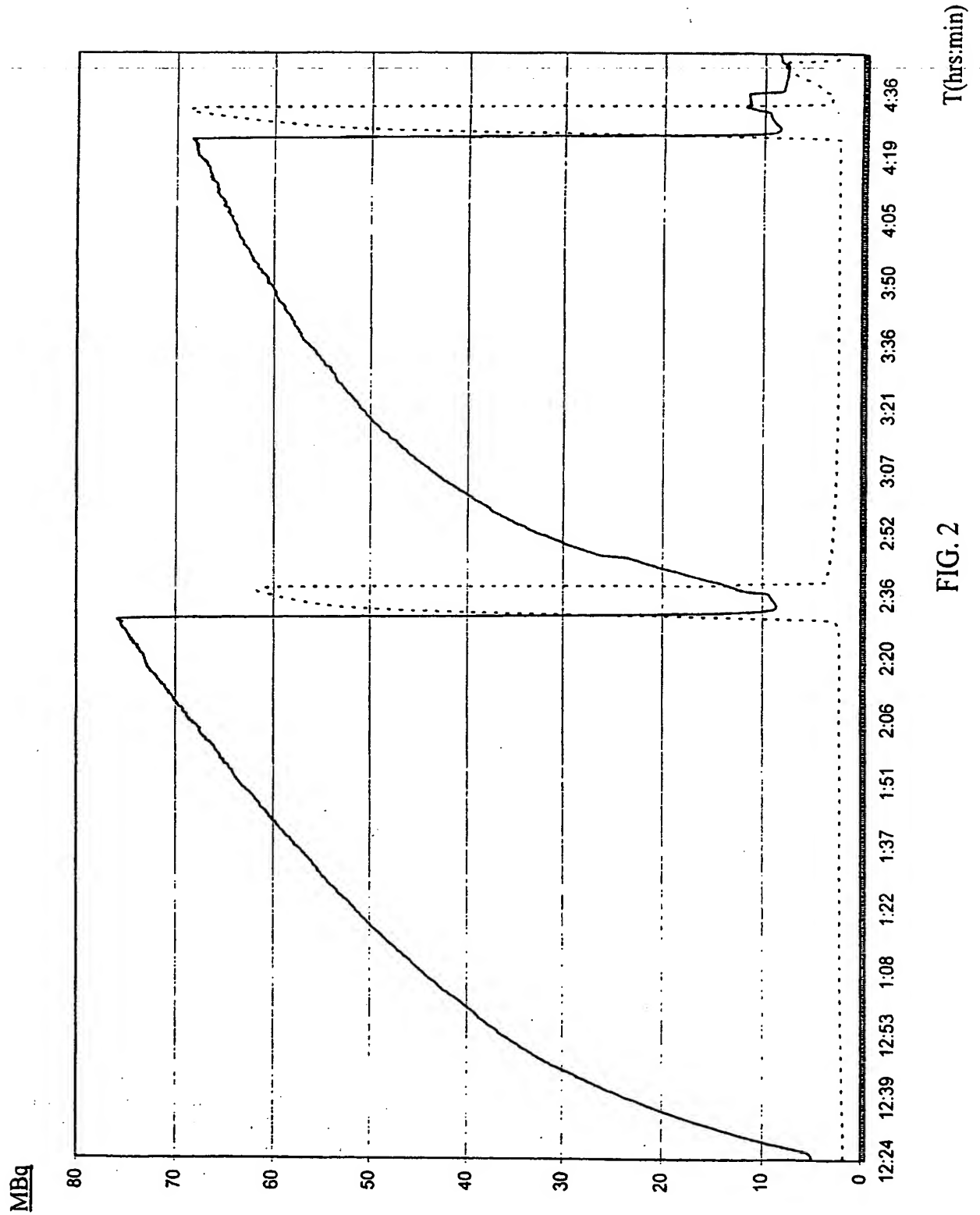


FIG. 2

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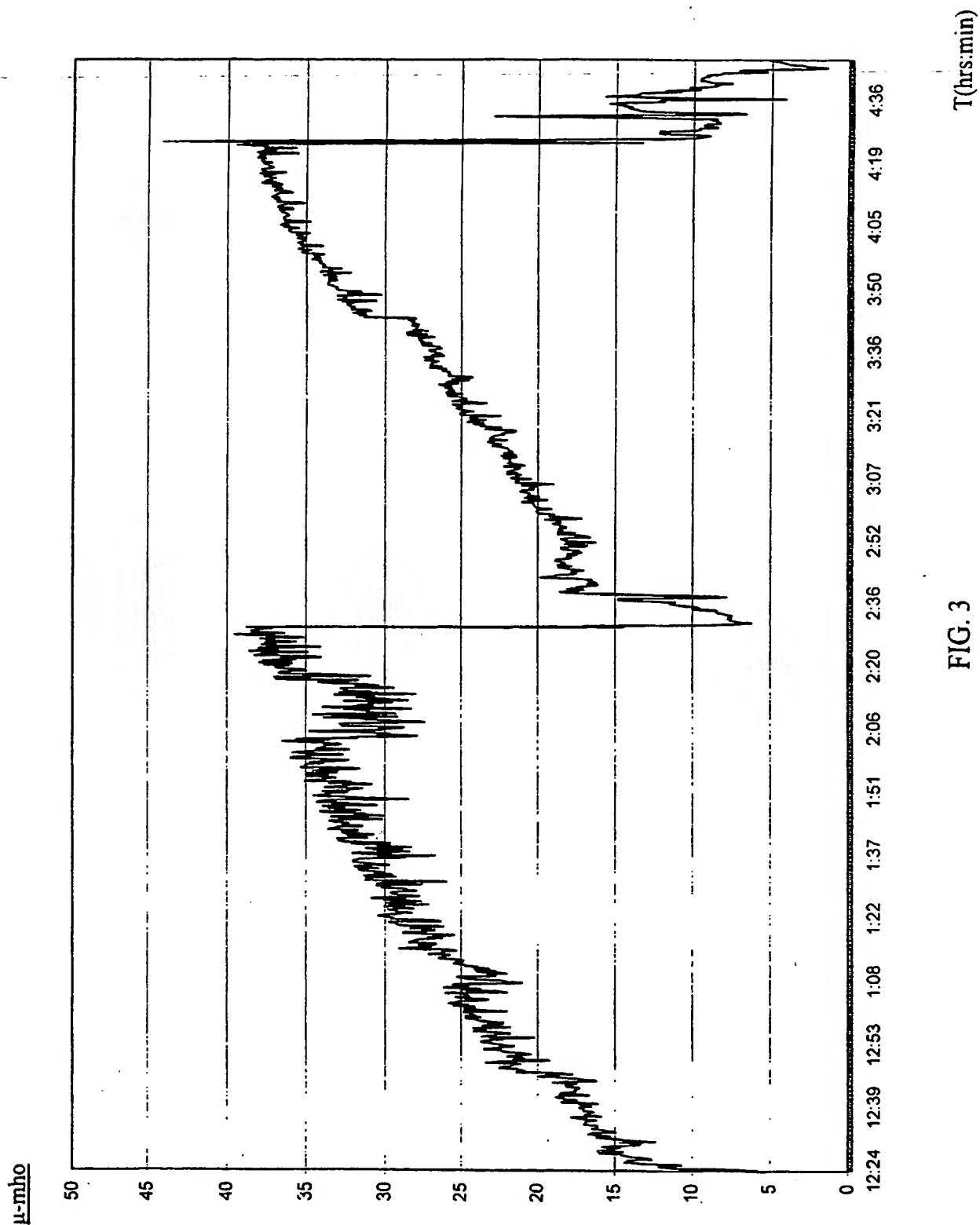


FIG. 3

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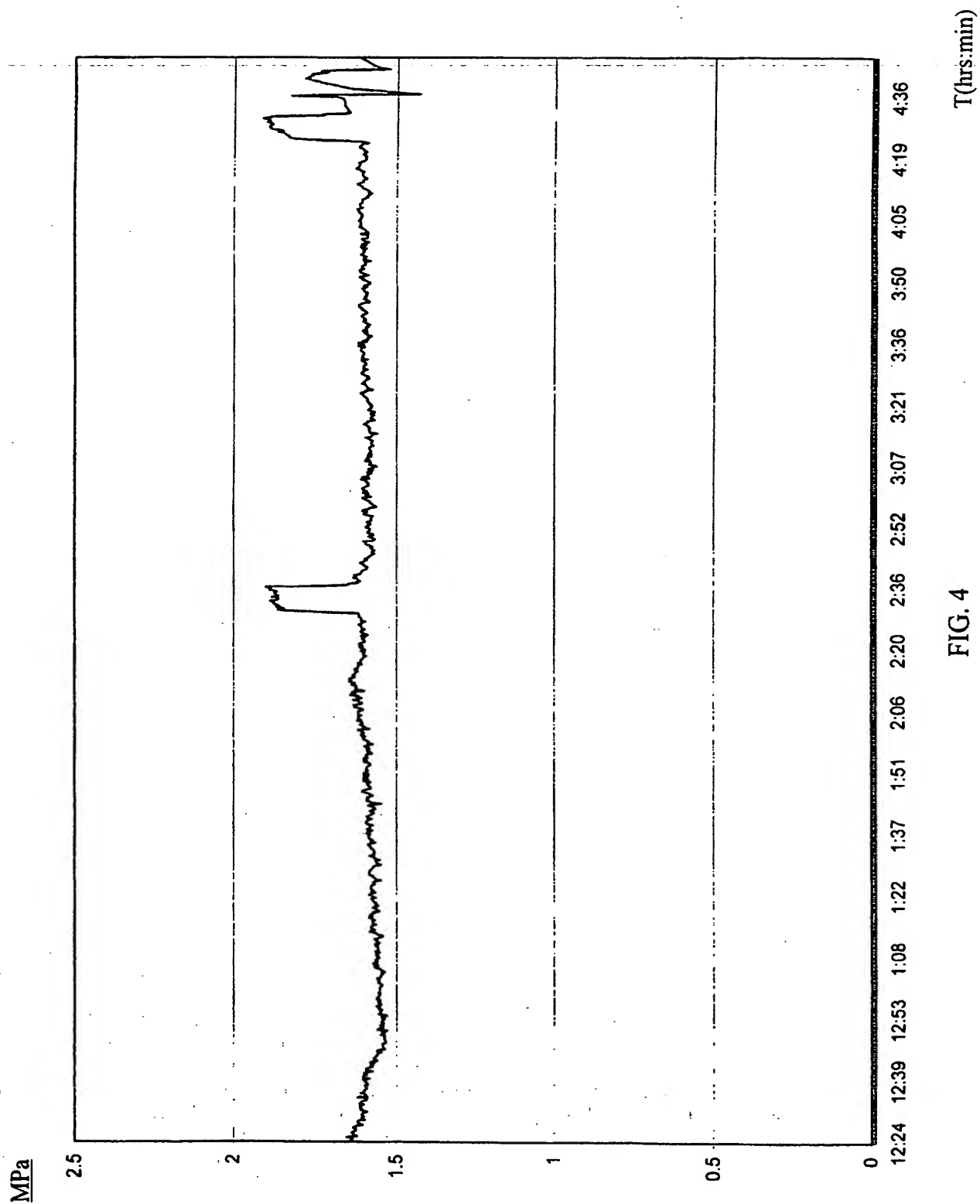


FIG. 4

## INTERNATIONAL SEARCH REPORT

National Application No

PCT/US 02/16017

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 7 G21G1/10

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**Minimum documentation searched (classification system followed by classification symbols)  
IPC 7 G21G

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, PAJ, INSPEC

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

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A	IWATA ET AL.: "18-F Fluoride production with a circulating 18-O water target" APPLIED RADIATION AND ISOTOPES, UK, vol. 38, no. 11, 1987, pages 979-984, XP001106132 the whole document	1,7
A	US 5 770 030 A (HAMACHER ET AL.) 23 June 1998 (1998-06-23) column 3, line 21 -column 4, line 10; figure 1	1,7
A	US 5 917 874 A (SCHLYER ET AL.) 29 June 1999 (1999-06-29) the whole document	1,7
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## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

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A	EP 0 949 632 A (NIHON MEDI-PHYSICS CO., LTD.) 13 October 1999 (1999-10-13) the whole document ---	1,7
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